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# Deletion of the signalling molecule synthase ScbA has pleiotropic effects on secondary metabolite biosynthesis, morphological differentiation and primary metabolism in *Streptomyces coelicolor* A3(2)

Davide D'Alia,<sup>1</sup> Daniela Eggle,<sup>2†</sup> Kay Nieselt,<sup>2</sup> Wei-Shou Hu,<sup>3</sup> Rainer Breitling<sup>4</sup> and Eriko Takano<sup>1\*</sup>

<sup>1</sup>Department of Microbial Physiology, Groningen Biomolecular Sciences and Biotechnology Institute, University of Groningen, Haren, the Netherlands.

<sup>2</sup>Center for Bioinformatics Tübingen, Department of Information and Cognitive Sciences, University of Tübingen, Tübingen, Germany.

<sup>3</sup>Department of Chemical Engineering and Material Science, University of Minnesota, Minneapolis, MN, USA.

<sup>4</sup>Institute of Molecular, Cell and Systems Biology, College of Medical, Veterinary and Life Sciences, University of Glasgow, Glasgow, UK.

## Summary

**Streptomyces** have high biotechnological relevance as producers of diverse metabolites widely used in medical and agricultural applications. The biosynthesis of these metabolites is controlled by signalling molecules,  $\gamma$ -butyrolactones, that act as bacterial hormones. In *Streptomyces coelicolor*, a group of signalling molecules called SCBs (*S. coelicolor* butanolides) regulates production of the pigmented antibiotics coelicolor polyketide (CPK), actinorhodin and undecylprodigiosin. The  $\gamma$ -butyrolactone synthase ScbA is responsible for the biosynthesis of SCBs. Here we show the results of a genome-wide transcriptome analysis of a *scbA* deletion mutant prior to and during the transition to antibiotic production. We report a strong perturbation in the expression of three pigmented antibiotic clusters in the mutant throughout the growth curve, thus providing a molecular explanation for the antibiotic phenotype observed previously. Our study also revealed, for the first time, that the secondary

metabolite cluster responsible for synthesis of the siderophore desferrioxamine is under the control of SCB signalling. Moreover, expression of the genes encoding enzymes for primary metabolism pathways, which supply antibiotic precursors and genes for morphological differentiation, was found shifted earlier in time in the mutant. In conclusion, our time series analysis demonstrates new details of the regulatory effects of the  $\gamma$ -butyrolactone system in *Streptomyces*.

## Introduction

Signalling molecules in bacteria play key functions in the intraspecies regulation of processes like differentiation, antibiotic production, biofilm formation and pathogenicity (Camilli and Bassler, 2006), and also contribute to inter-species communication (Subramoni and Venturi, 2009). The molecules act at a very low concentration, and therefore are often referred to as microbial hormones. The discovery of these molecules has opened a new research area and has important applications in biotechnology, for instance related to biofilm formation (Landini *et al.*, 2010).

*Streptomyces* are biotechnologically important bacteria and well known for their ability to produce numerous secondary metabolites, including anticancer agents, immunosuppressants and most importantly antibiotics, of which they are the major producer. The best-characterized signalling molecule in *Streptomyces* is A-factor, the  $\gamma$ -butyrolactone of *Streptomyces griseus* (Khokhlov *et al.*, 1967). The key enzyme in the synthesis of A-factor has been shown by *in vitro* synthesis to be AfsA (Kato *et al.*, 2007). Other  $\gamma$ -butyrolactones that have been characterized are the virginiae butanolides (VBs) from *Streptomyces virginiae*. The *afsA* homologue in *S. virginiae* is *barX*; its deletion causes a deficiency in VBs and virginiamycin production (Lee *et al.*, 2010). It has recently been reported that addition of VBs to a *barX* null mutant restores the wild-type phenotype with respect to virginiamycin production (Lee *et al.*, 2010), confirming its enzymatic activity. Also, BarX stabilizes interaction between BarA and the *barB* promoter (Kawachi *et al.*,

Received 23 June, 2010; accepted 12 October, 2010. \*For correspondence. E-mail e.takano@rug.nl; Tel. (+31) 50 3632143; Fax (+31) 50 3632154. †Present address: Pharma Research Scientific Informatics, In Silico Sciences- Bioinformatics, Roche Diagnostics GmbH, Pharma Research Penzberg, Penzberg, Germany.

2000), suggesting a regulatory function beside the enzymatic role.

In *Streptomyces coelicolor* A3(2), three  $\gamma$ -butyrolactones have been determined as signalling molecules that elicit antibiotic production (Takano *et al.*, 2000; Hsiao *et al.*, 2009), of which the most abundant is the *S. coelicolor* butanolide 1 (SCB1). SCBs are known to regulate at least three antibiotic biosynthetic clusters: *act*, for the synthesis of the blue-pigmented actinorhodin (Act); *red*, for the synthesis of the red-pigmented undecylprodigiosin (Red); and the recently characterized *cpk* for the synthesis of the yellow pigment  $\gamma$ CPK and the antibiotic abCPK (Takano *et al.*, 2005; Gottelt *et al.*, 2010).

One of the strategies to increase productivity of a specific antibiotic in streptomycetes is the removal of  $\gamma$ -butyrolactones receptor proteins (Martin and Liras, 2010). For example, deletion of *tylP* in *Streptomyces fradiae* (Cundliffe, 2008) and *brp* in *Streptomyces clavuligerus* (Santamarta *et al.*, 2005) have been reported to increase, between two- and threefold, the production of tylosin, and clavulanic acid and cephamycin C respectively. Deleting the  $\gamma$ -butyrolactones receptor *scbR* in *S. coelicolor* does not trigger overproduction of the Red and Act antibiotics, but rather delays their synthesis (Takano *et al.*, 2001).

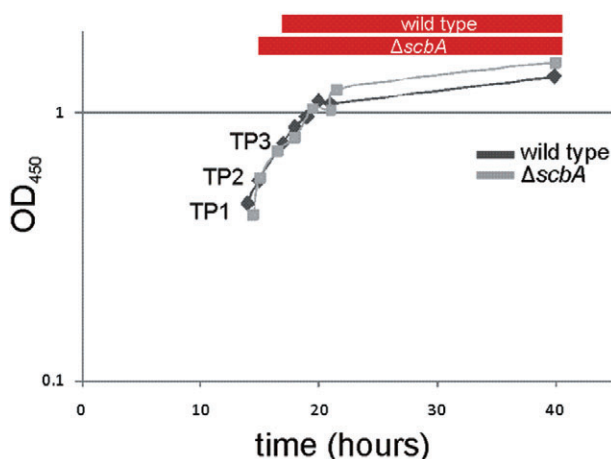
The AfsA homologue in *S. coelicolor*, ScbA, is involved in SCB production which occurs during transition and stationary phase (Takano *et al.*, 2001). It has homology to fatty acid synthases, and mutation of two predicted enzymatically active sites leads to the loss of SCBs (Hsiao *et al.*, 2007). ScbR is the main butanolide receptor in *S. coelicolor* and acts as transcriptional regulator (Takano *et al.*, 2001). In the absence of the signalling molecules, ScbR binds to the promoter of its own gene (site R) and of *cpkO* (site OA), the main activator of the *cpk* gene cluster (Takano *et al.*, 2005), repressing their expression. Also, ScbR binds the promoter of *scbA* (site A); however, with an opposite effect: its presence is necessary for *scbA* induction (Takano *et al.*, 2001). *scbA* is divergent from *scbR*, and its deletion impairs production of SCBs, and causes an earlier and enhanced Red and Act production (Takano *et al.*, 2001). *scbA* and *scbR* expression is absent in the *S. coelicolor* *scbA* deletion mutant M751; addition of SCB1 restores *scbR* transcription, but not that of *scbA*. These data taken together suggest a possible regulatory function of ScbA, at least for its own expression (Takano *et al.*, 2001). To analyse the role of ScbA and the butanolide system more extensively, gene expression in a *scbA* deletion mutant (M751), just before and during the onset of antibiotic production, was compared with the wild-type M145 using whole-genome microarrays and quantitative RT-PCR (qRT-PCR) analyses.

## Results

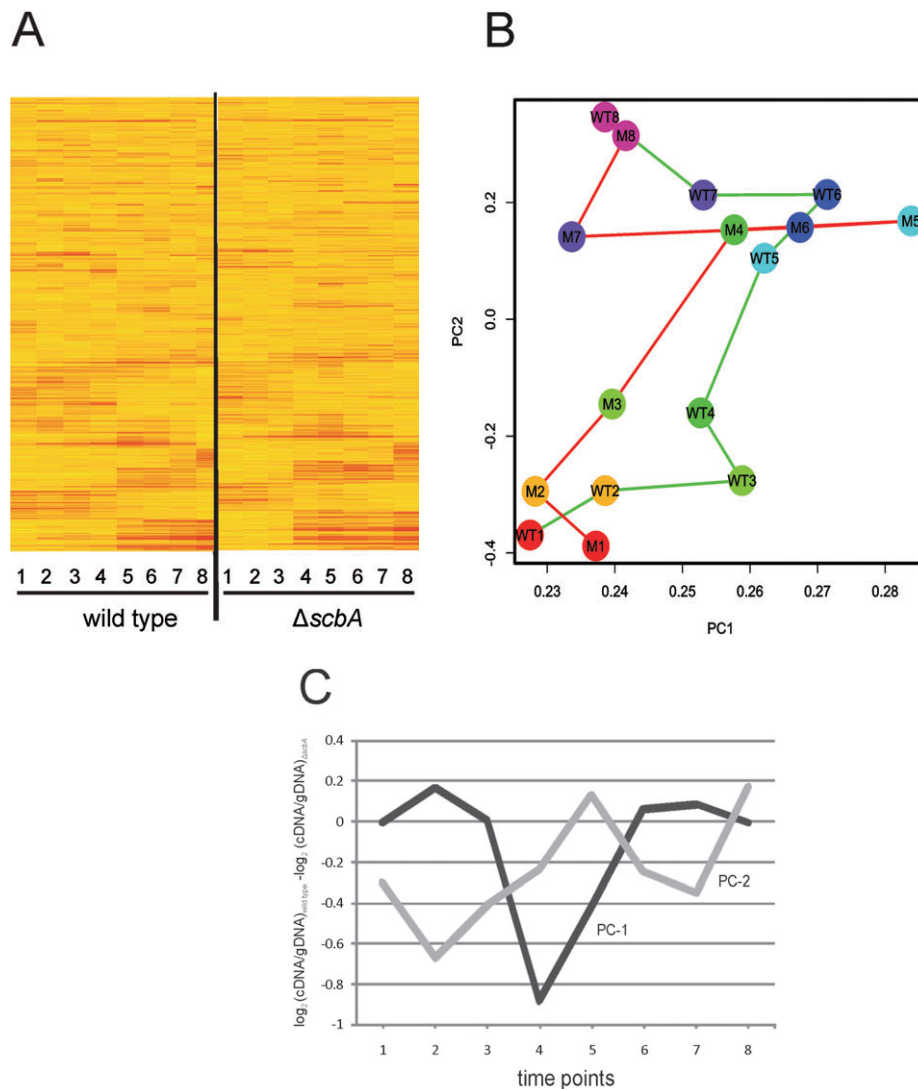
### Microarray analysis of differentially expressed genes in the $\gamma$ -butyrolactone non-producer mutant versus the wild type

Antibiotic production in the *scbA* deletion mutant (M751) occurs earlier and leads to a 10-fold higher final yield of both Act and Red than that of the wild-type (M145) strain (Takano *et al.*, 2001). To understand the possible regulatory role of ScbA and how ScbA influences the production of antibiotics, the genome-wide time-series expression profiles of *S. coelicolor* wild-type M145 and M751 were compared by microarray analysis. The two strains showed the same growth profile in SMM liquid medium (Fig. 1). While the wild type started producing the red-pigmented undecylprodigiosin at time point 3 (TP3) (red bar in Fig. 1), at the switch from exponential to transition phase, followed by the blue pigmented actinorhodin, production of Red in the mutant occurred during exponential phase at TP2 (red bar in Fig. 1). RNA was isolated from eight time points during the growth curve and cDNA retrotranscribed and hybridized to a PCR-based microarray comprising 7629 genes. The raw expression data are available at the following website: <http://www-ps.informatik.uni-tuebingen.de/coelicolor>.

The overall expression trend of the majority of the genes was similar between the two strains (Fig. 2A). This was expected given that the two strains show a similar growth profile (Fig. 1). The expression profiles at all time points were analysed using principal component analysis (PCA). In the plot of the first two principal components, one can see that the two strains follow very



**Fig. 1.** Growth curves of the wild-type and the  $\Delta scbA$  mutant. Cell growth in liquid SMM medium of the wild type (dark line, diamond) and the  $\Delta scbA$  mutant (light line, square) was monitored in time, measuring the absorbance of the culture at OD<sub>450</sub>. Undecylprodigiosin production started at the beginning of the transition phase in the wild type (red bar) and at mid-exponential phase in the  $\Delta scbA$  mutant (red bar).



**Fig. 2.** Identification of perturbed genes in the  $\Delta scbA$  mutant using principal component analysis (PCA).

A. Microarray analysis of global transcription profile of the wild type (left panel) and the  $\Delta scbA$  mutant (right panel), sorted by hierarchical clustering. Intensity of the colour (from yellow to red) represents intensity of the expression for each gene.

B. The gene expression values in wild type (green line) and the  $\Delta scbA$  mutant (red line) were analysed by PCA, and the values of principal component 1 are plotted versus principal component 2. Numbers in the circles indicate the TPs.

C. The two main patterns of the difference expression profile between the wild type and the  $\Delta scbA$  mutant are shown, based on the loadings of PC-1 (dark line) and PC-2 (light line) in PCA of the expression difference profiles.

similar curves, but one can also notice a shift occurring around TP3, just as reflected in the antibiotic production (Fig. 2B).

#### Differential expression analysis at time point 1

To elucidate the cause of the earlier production of antibiotics in M751, we characterized in detail the expression profile of genes at early exponential phase (time point 1), which is prior to the production of antibiotics in M751. Differentially expressed genes were calculated as described in *Experimental procedures*. Fifty-six genes

were selected based on these criteria, which were either upregulated at least 1.2-fold or downregulated at least 0.85-fold in M751 compared with M145 (Table S1). qRT-PCR was carried out for 34 of the 56 differentially expressed genes. *hrdB* was chosen as the endogenous control. Eighteen out of the 25 upregulated and one out of the nine downregulated genes were confirmed (Table 1). Of the genes whose expression was confirmed by qRT-PCR, the largest group was encoding hypothetical proteins, the second largest consisting of genes with house keeping functions. SCO0982, encoding the isocitrate lyase *aceA*, was the second most overexpressed

**Table 1.** Differentially expressed genes at TP1 by qRT-PCR.

Gene	Annotation and common name	Functional annotation	Fold change
SCO0607	<b>Hypothetical lipoprotein</b>	<b>Periplasmic/exported/lipoproteins</b>	<b>6.53</b>
SCO0982	<b>Isocitrate lyase <i>aceA</i></b>	<b>Central intermediary metabolisms</b>	<b>2.23</b>
SCO2476	Putative dehydrogenase/reductase (putative secreted protein)	Not classified (included putative assignments)	2.23
SCO3752	Putative ABC transporter ATP-binding protein	Cell process	2.02
SCO4295	<b>Cold shock protein <i>scoF4</i></b>	<b>Cell process</b>	<b>1.93</b>
SCO4425	<b>Sigma-like protein <i>afsS</i></b>	<b>Other regulation</b>	<b>1.75</b>
SCO3225	Two-component sensor kinase <i>absA</i>	Secondary metabolism	1.81
SCO6577	<b>Conserved hypothetical protein</b>	<b>Hypothetical protein</b>	<b>1.65</b>
SCO5865	Hypothetical protein	Hypothetical protein	1.65
SCO3538	<b>Hypothetical protein</b>	<b>Hypothetical protein</b>	<b>1.53</b>
SCO6624	<b>Putative membrane protein</b>	<b>Hypothetical protein</b>	<b>1.52</b>
SCO3893	<b>Hypothetical protein</b>	<b>Hypothetical protein</b>	<b>1.51</b>
SCO7476	<b>Putative secreted protein</b>	<b>Periplasmic/exported/lipoproteins</b>	<b>1.50</b>
SCO3133	<b>Putative MarR-family regulator</b>	<b>Defined family regulators</b>	<b>1.48</b>
SCO2966	<b>Small protein B homologue <i>smpB</i></b>	<b>Macromolecule metabolism</b>	<b>1.47</b>
SCO4173	Hypothetical protein	Hypothetical protein	1.46
SCO2261	<b>Putative membrane protein</b>	<b>Periplasmic/exported/lipoproteins</b>	<b>1.42</b>
SCO2195	<b>Hypothetical protein</b>	<b>Hypothetical protein</b>	<b>1.39</b>
SCO1152	<b>Putative helicase</b>	<b>Macromolecule metabolism</b>	<b>1.36</b>
SCO5725	<b>Hypothetical protein</b>	<b>Hypothetical protein</b>	<b>1.32</b>
SCO0981	Putative DNA-binding protein	Other regulation	1.27
SCO4296	<b>Chaperonin 2 <i>groEL2</i></b>	<b>Cell process</b>	<b>1.24</b>
SCO6259	<b>Probable ABC sugar transport ATP-binding protein</b>	<b>Cell process</b>	<b>1.24</b>
SCO3669	<b>Molecular chaperone <i>dnaJ</i></b>	<b>Cell process</b>	<b>1.19</b>
SCO2140	Putative transcriptional regulator	Defined family regulators	1.11
SCO6336	Hypothetical protein	Hypothetical protein	1.10
SCO5998	Putative bifunctional protein (putative MurA2)	Periplasmic/exported/lipoproteins	1.09
SCO2950	DNA-binding protein Hu <i>hs1</i>	Other regulation	1.07
SCO5494	Putative DNA ligase <i>ligA</i>	Macromolecule metabolism	0.88
SCO4727	30S ribosomal protein S13 <i>rpsM</i>	Ribosome	0.81
SCO4709	50S ribosomal protein L16 <i>rplP</i>	Ribosome	0.80
SCO6099	<b>Adenylylsulfate kinase <i>cysC</i></b>	<b>Central intermediary metabolisms</b>	<b>0.76</b>
SCO2198	Glutamine synthetase I <i>glnA</i>	Amino acid biosynthesis	0.76
SCO5368	ATP synthase C chain <i>atpE</i>	Energy metabolism; carbons	0.66

Thirty-four genes differentially regulated in M751 compared with M145 at TP1 selected from the microarray were analysed by qRT-PCR. Their ID, annotation and functional annotation are indicated in the table. Also the fold change of expression in M751 compared with M145 is reported in the last column. In bold, the genes whose differential expression have been confirmed by microarray and qRT-PCR, independently. The genes are sorted by fold change.

gene. This gene is essential for growth on acetate as sole carbon source (Akopiants *et al.*, 2006), and its maximum activity has been observed during cessation of growth in *S. clavuligerus* (Soh *et al.*, 2001). A gene encoding a cold-shock protein, *scoF4*, was the third most overexpressed. Similarly, three other genes upregulated in the mutant are related to stress response: the transfer-messenger RNA cofactor *smpB*, and the molecular chaperones *dnaJ* and *groEL2*. *afsS*, a sigma-like protein, which is a pleiotropic regulator for antibiotics production, was identified as the fourth most overexpressed gene. The only gene that was found downregulated independently by microarray and qRT-PCR was *cysC*, encoding an adenylylsulfate kinase, part of the biosynthetic pathway for the amino acid cysteine. Some genes whose expression was downregulated in the microarray analysis were in fact upregulated when analysed by qRT-PCR. *absA1* was identified as such and it is interesting to note that it is part of the two-component system which was

shown to regulate Act and Red production (Anderson *et al.*, 2001).

#### Global expression analysis of the growth curve

Moreover, to explore the global effects on the transcriptome during the entire growth curve, we calculated average log<sub>2</sub> expression ratios of mutant versus wild-type expression at each time point. The differential expression profiles at all time points were analysed using PCA to identify major patterns in differential expression across the growth curve. The two major patterns are shown in Fig. 2C. Principal component 1 (PC-1), which accounted for 24% of the variance in the difference profiles, contained genes which displayed a tendency to be lower expressed in M145 at time points 4 and 5 compared with M751 (Fig. 2C). Principal component 2 (PC-2), which accounted for 19% of the variance, showed that some genes were lower expressed in the wild type at all time



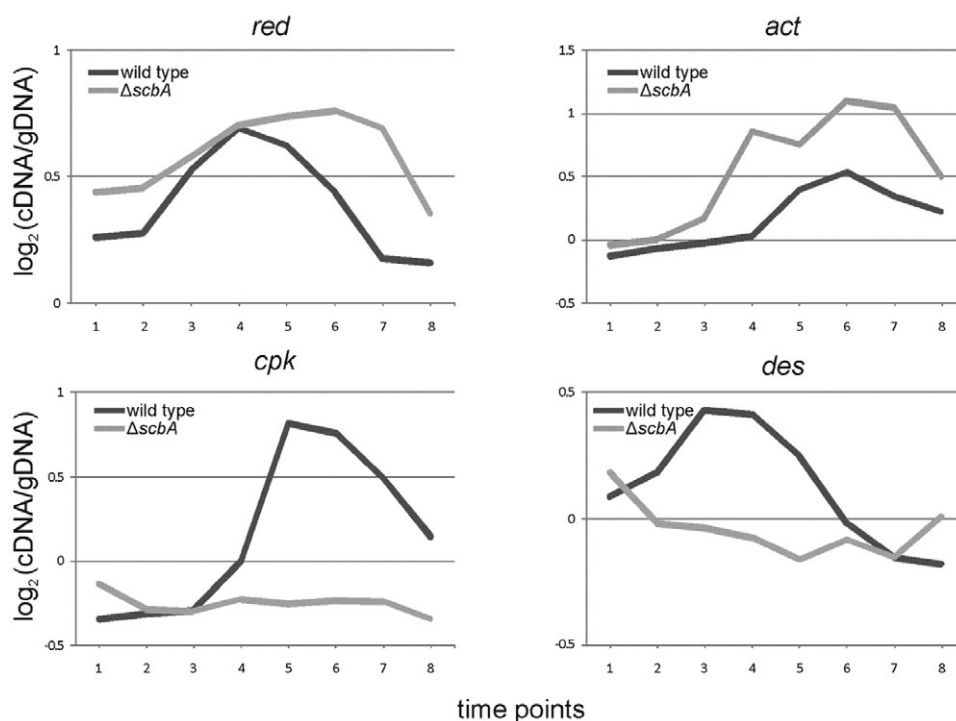
points before and after transition phase (TP5) compared with the mutant (Fig. 2C). Among the genes with the most extreme values along either PC-1 or PC-2, two main groups were identified with characteristic differential expression profiles. The first group comprised mostly genes involved in secondary metabolism that were upregulated (or silenced) in the mutant from exponential phase (TP1 or TP2) until late stationary phase (TP8). The second group consists of those genes with expression profiles that are alike but shifted by one time point earlier in the mutant compared with the wild type; this group contained mostly genes involved in primary metabolism and morphological differentiation, as well as the conservation 9 (discussed below).

*Group I: gene clusters up- or down-regulated throughout the time series*

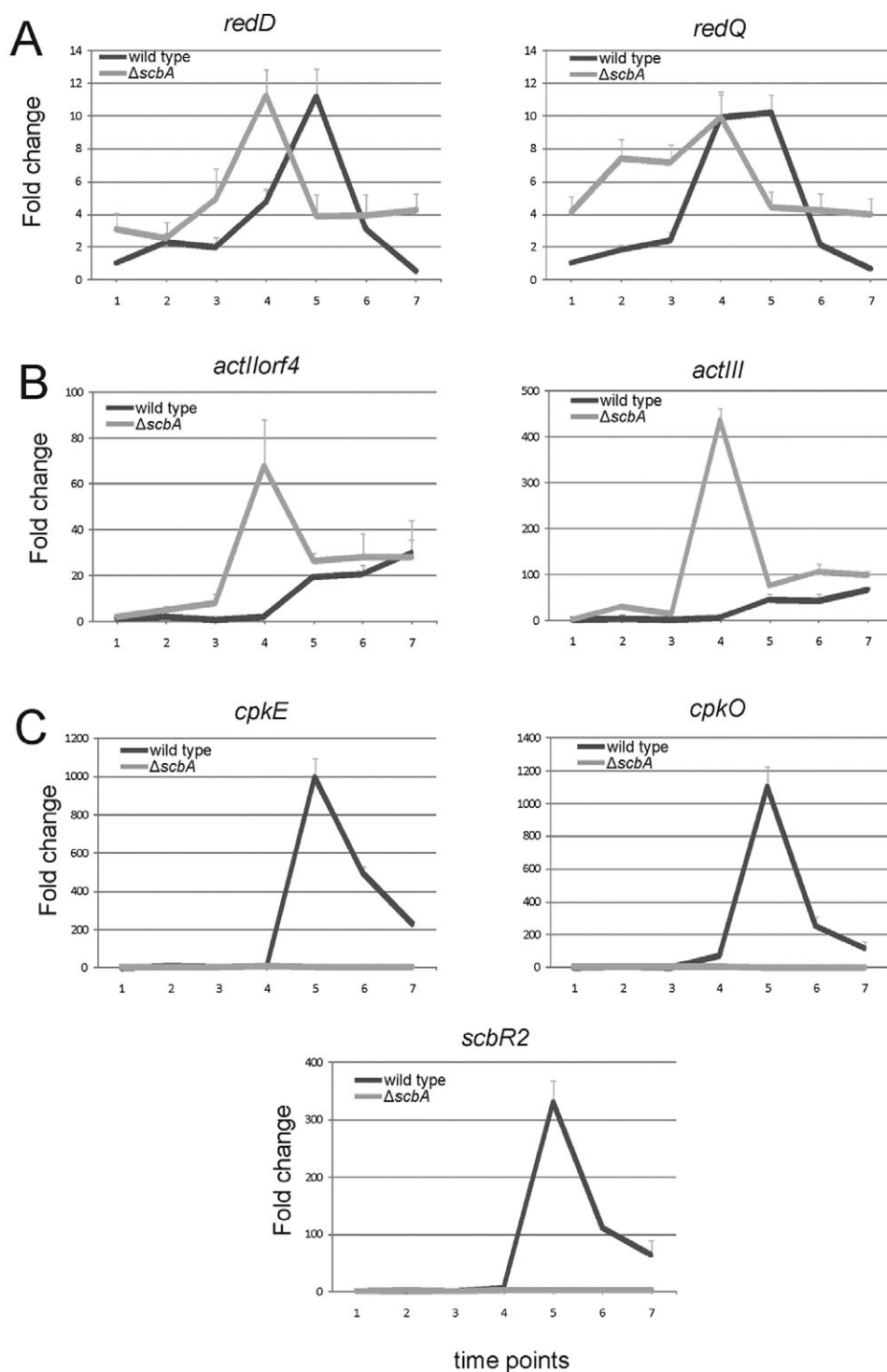
This group mainly consists of secondary metabolism gene clusters. Consistent with the observed earlier and more intense production of undecylprodigiosin, the microarray analysis revealed that the average of genes of the *red* biosynthetic cluster (SCO5877–5898) were expressed at higher values all along the growth curve and decreased in expression only very late at TP8 in the mutant, while in the wild-type transcription peaked at TP4 and TP5 (transition

phase) and decreased immediately afterwards (Fig. 3). In addition to undecylprodigiosin, *scbA* mutant M751 overproduced the blue-pigmented actinorhodin, also consistent with an earlier report (Takano *et al.*, 2001). The expression profile of its biosynthetic cluster *act* (SCO5071–5091) peaked at TP5 and TP6 (transition and early stationary phase) in the wild type, but was already almost twofold higher at TP4 in the mutant (Fig. 3). Earlier expression of *redD* and *redQ* (the main activator and the acyl carrier protein in the *red* cluster respectively), *actII-orf4* and *actIII* (the main activator and the ketoacyl reductase in the *act* cluster respectively) was confirmed by qRT-PCR (Fig. 4A and B).

Most dramatically, the *cpk* cluster (SCO6268–6288) was silenced in M751 throughout the entire growth curve (Fig. 3), while in the wild type it showed a sharp overexpression from TP4 onwards, coinciding with the expression peak of *scbA* and *scbR* (Fig. S1). Also, by qRT-PCR analysis, the expression of three genes in the cluster, namely *cpkE* (homologue to epoxide hydrolase) and the two regulators *cpkO* and *scbR2* [homologue to *scbR* and negative regulator of the cluster (Gottelt *et al.*, 2010)], was confirmed to be the same as the microarray analysis (Fig. 4C). Interestingly, the direct neighbours of the *cpk* cluster, SCO6265 (*scbR*) and SCO6267 (*scbC*) (Fig. S1), were also drastically downregulated. SCO6265 encodes



**Fig. 3.** Expression profiles of secondary metabolism genes in the wild type and the  $\Delta scbA$  mutant. The average expression profiles of the secondary metabolite biosynthetic clusters *red* (SCO5877–5898), *act* (SCO5071–5091), *cpk* (SCO6268–6288) and *des* (SCO2782–2785) during the time-course are plotted for the wild type (dark line) and the  $\Delta scbA$  mutant (light line). A list of the genes and their expression data is included in Table S3.



**Fig. 4.** Quantitative RT-PCR for selected genes. Expression profiles of genes of the *red* cluster [*redD* (SCO5877) and *redQ* (SCO5887)] (A), the *act* cluster [*actII-orf4* (SCO5085) and *actIII* (SCO5086)] (B), the *cpk* cluster [*cpkE* (SCO6277), *cpkO* (SCO6280) and *scbR2* (SCO6286)] (C) are plotted for the wild type (dark line) and the  $\Delta scbA$  mutant (light line). The expression for each gene is compared with the value in M145 at TP1, which is set as 1. Standard deviation at each time point is denoted in a thin light line.

the  $\gamma$ -butyrolactones receptor ScbR while SCO6267 encodes the  $\gamma$ -butyrolactones synthase ScbC (Nieselt *et al.*, 2010) and has 76% identity in amino acid sequence to BrpA, an oxidoreductase involved in reduction of the butanolide during A-factor biosynthesis in *S. griseus* (Kato *et al.*, 2007). The desferrioxamine E biosynthetic cluster *des* (SCO2782–SCO2785) has a reverse expression profile, compared with the *red* and *act* clusters, and was strongly downregulated from TP2 onwards in M751 (Fig. 3). Desferrioxamine E is the main siderophore in *S. coelicolor* (Barona-Gomez *et al.*, 2004).

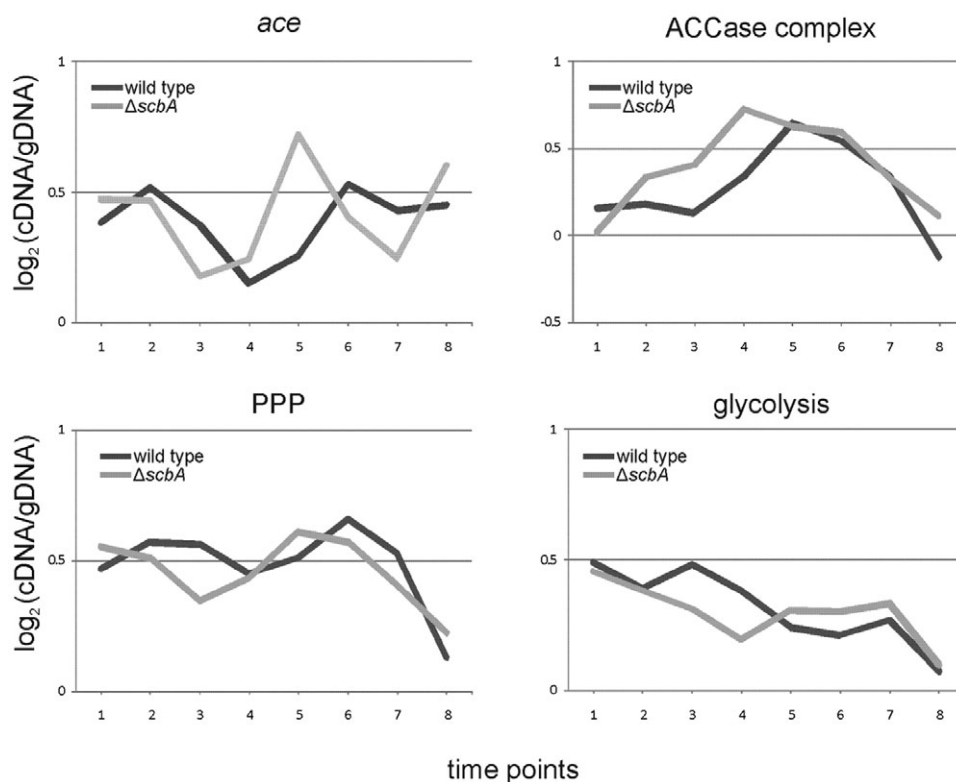
*Group II: gene clusters shifted by one time point earlier in the mutant compared with the wild type*

Our microarray analysis showed that the expression profile of *ace* (for conversion of pyruvate to acetyl-CoA), the ACCase complex (from acetyl-CoA to malonyl-CoA) and the PPP (pentose phosphate pathway) genes is shifted one time point earlier in M751 compared with the wild type (Fig. 5). This pattern is weak for individual genes, but highly consistent across the set of genes involved. Similarly, although to a somewhat less extent, the gene expression of the glycolysis pathway was also

shifted (Fig. 5). *Streptomyces coelicolor* has been shown to differentiate in liquid media (Manteca *et al.*, 2008). A set of genes involved in morphological differentiation were found expressed and appeared to be shifted to one time point earlier in M751 (Fig. 6). These genes are *ramS*, encoding for SapB (Kodani *et al.*, 2004), *nepA* (de Jong *et al.*, 2009), *chpH* (Claessen *et al.*, 2003), *bldG* and *sigK* (Mao *et al.*, 2009a; Parashar *et al.*, 2009). Another cluster showing a shifted expression profile to one time point earlier in the mutant compared with the wild type was the conservon 9 (Fig. 6). Conservons (cvns) are operons identified in *S. coelicolor* (Bentley *et al.*, 2002) and conserved specifically in species belonging to *Actinomycetales* (Komatsu *et al.*, 2006).

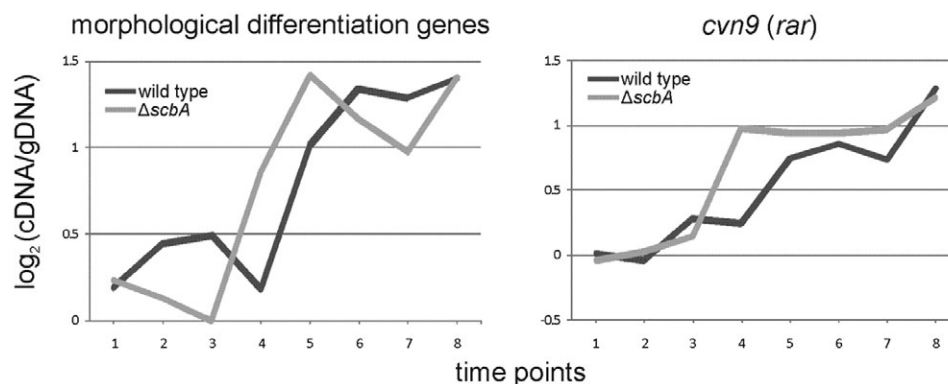
*Putative non-coding RNAs are differentially regulated in M751*

Our group has recently predicted the presence of small non-coding RNAs (ncRNAs) in inter- and intragenic loci in the *S. coelicolor* genome and experimentally confirmed the presence of 13 antisense ncRNAs (D'Alia *et al.*, 2009). Because of technical reasons (the microarray in this study is PCR-based) it was not possible to separately



**Fig. 5.** Expression profiles of primary metabolism genes in M145 and the  $\Delta scbA$  mutant. The average of genes involved in conversion of pyruvate to acetyl-CoA (*ace*) and from acetyl-CoA to malonyl-CoA (ACCase complex) and the average expression profiles of the primary metabolism genes for the PPP and glycolysis pathways during the time-course are plotted for M145 (dark line) and the  $\Delta scbA$  mutant (light line). A list of the genes and their expression data is included in the Table S3.





**Fig. 6.** Expression profiles of morphological differentiation genes and conservon 9. The average expression profiles of a set of morphological differentiation genes and the conservon 9 [*cvn9 (rar)*] are plotted for M145 (dark line) and the  $\Delta scbA$  mutant (light line). A list of the genes and their expression data are included in Table S3.

analyse the messenger and antisense transcripts. Nevertheless, 11 short coding sequences (maximum length 300 bp), which we predicted as ncRNA and which are annotated as encoding unknown or hypothetical proteins, were strongly upregulated in the  $\gamma$ -butyrolactone-impaired mutant at TP4 (Fig. 7).

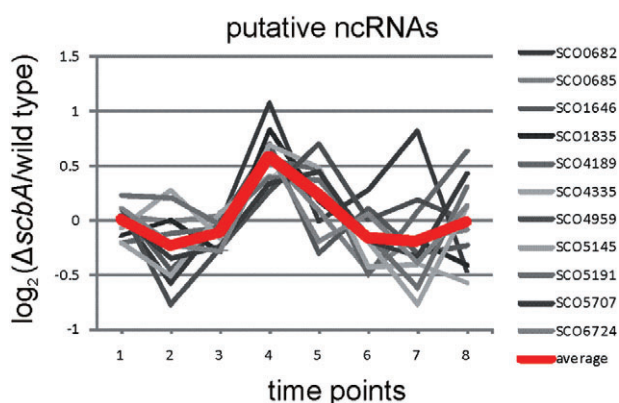
## Discussion

The microarray and qRT-PCR analyses reported in this study showed that the earlier and more intense production of Act and Red in the  $\gamma$ -butyrolactone signalling mutant M751 is the direct effect of changes in the expression of their biosynthetic clusters. This observation points at the involvement of the butanolide system in regulating the *act* and *red* clusters at a transcriptional level, perhaps via an intermediate regulator. Furthermore, we could show that the *cpk* cluster was silenced throughout the growth in the

mutant. These data agree with the reported abolished expression of *cpkO*, the pathway-specific activator of the *cpk* cluster in the *scbA* mutant (Takano *et al.*, 2005). This explains the previously reported discrepancy in the  $\gamma$ -butyrolactone synthase mutant antibiotic production phenotype observed in *S. coelicolor* compared with that in *S. griseus* and *S. virginiae* (Hara and Beppu, 1982; Kawachi *et al.*, 2000). In a fashion similar to the systems in *S. griseus* and *S. virginiae*, where deleting the signalling molecules synthase impairs streptomycin and virginiamycin production, deleting the butanolides synthase *ScbA* impairs *cpk* expression.

Another secondary metabolite cluster drastically affected by deleting *scbA* was the siderophore desferrinoxamine E biosynthesis. In contrast, the biosynthetic clusters of the siderophore coelichelin (Lautru *et al.*, 2005) and the siderophore-like coelibactin (Hesketh *et al.*, 2009) did not show any remarkable difference in expression between the two strains (data not shown). Taken together, this would presumably lead to a decrease in intracellular iron levels. Interestingly, it has been shown that iron limitation enhances intracellular production and export of actinorhodin in *S. coelicolor* (Coisne *et al.*, 1999). To our knowledge, this is the first report of a siderophore whose control involves the  $\gamma$ -butyrolactone system.

In a similar context, it has been reported that blocking the A-factor cascade in *S. griseus* results in strong changes in secondary metabolism (Ohnishi *et al.*, 2008; Hara *et al.*, 2009). The two cited reports showed that in addition to the streptomycin biosynthesis genes, several other genes for secondary metabolites were induced by the A-factor system: three putative non-ribosomal peptide synthetase (NRPS) clusters, a type I polyketide synthase (PKS), a lantibiotic and a putative melanin biosynthesis clusters, a putative terpene cyclase and two signalling molecules receptor homologues.



**Fig. 7.** Expression profiles of putative ncRNAs. The fold change between the  $\Delta scbA$  mutant and M145 of expression profiles of 11 genes (indicated at the side) overlapping putative ncRNAs are plotted. The average is emphasized by a thick red line. A list of the genes and their expression data is included in Table S3.

Another very important effect shown by the global gene expression analysis was the apparent earlier shift of the primary metabolism genes. This includes the genes for PPP and glycolysis pathways, and genes for the conversion of pyruvate to acetyl-CoA (*ace*) and malonyl-CoA (*acc*). This effect would have been impossible to discover in a more targeted analysis. These findings agree with the earlier production of antibiotics, suggesting that the deletion of *scbA* causes a shift of the primary metabolism pathways, which is required for Act and Red production. In particular, earlier and more intense expression of genes encoding the ACCase complex in M751 increases the amount of available malonyl-CoA, an essential precursor for Act and Red biosynthesis (Cerdano *et al.*, 2001; Beltran-Alvarez *et al.*, 2007). This is also in agreement with the observation that the peak in expression of the *ace* genes is at TP5 in the mutant, when also the *act* cluster is actively transcribed. In this phase, the cell needs more acetyl-CoA to sustain synthesis of malonyl-CoA. These findings agree with the reported increase of actinorhodin by increasing the levels of its precursor malonyl-CoA by overexpressing acetyl-CoA carboxylase *acc* (Ryu *et al.*, 2006). On the other hand, the earlier decrease in expression of the glycolysis genes in M751 could preserve glucose-6-P to be used in the PPP which is the main source of NADPH, the source of reducing power for secondary metabolism (Butler *et al.*, 2002).

A similar shift was observed in the expression of morphological differentiation genes and conservon 9. Our transcriptional analysis revealed that several genes involved in morphological differentiation were expressed in M145 and M751 grown in liquid. Recently, it has been reported that *S. coelicolor* expresses developmental genes in liquid cultures and that their expression increases in time until transition phase, and then is switched off (Nieselt *et al.*, 2010). Our analysis revealed an earlier expression of the morphological differentiation genes in M751. Earlier production of aerial hyphae or spores have not been reported in the *scbA* mutant (Takano *et al.*, 2001), and we also did not observe any difference in development on solid media between the wild type and the mutant. However, the earlier expression of these genes may cause a subtle shift in time of the morphological differentiation which we cannot readily observe under the conditions used. Further analysis will be needed to reveal a potential involvement of SCBs in differentiation, as has been observed for the  $\gamma$ -butyrolactones in *S. griseus* (Horinouchi, 2002), *S. ansochromogenes* (Li *et al.*, 2003) and *S. natalensis* (Lee *et al.*, 2005). A similar shift in expression of conservon 9 was also detected. Thirteen conservons have been identified in *S. coelicolor* (Bentley *et al.*, 2002), and they typically consist of two genes with unknown function sandwiched between a sensor histidine kinase and an

ATP/GTP-binding protein. In four cases (*cvn9–12*) a cytochrome P450 is also found in the gene cluster. Disruptions of *cvnA9*, a histidine kinase homologue, and *cvnD9*, homologous to ATP/GTP-binding protein, have been reported to lead to conditional overproduction of actinorhodin (Komatsu *et al.*, 2003; Komatsu *et al.*, 2006). Interestingly, the increase in expression of *cvn9* overlaps with the expression peak of the *act* cluster, both in M145 (TP5) and M751 (TP4).

Recently, our group has shown the presence of 13 intragenic antisense ncRNAs and that at least two of them, namely *cnc3669.2* and *cnc3893.2*, were not expressed in the wild type but were detected in the two  $\gamma$ -butyrolactone deficient mutants M751 and M752 (*scbR* deletion mutant). We suggested a role of the signalling molecules in regulating the expression of these small transcripts (D'Alia *et al.*, 2009). Eleven CDS encoding hypothetical proteins which overlapping with our predicted ncRNAs have been found upregulated in M751 at transition phase (TP4). Among the 11 CDS, SCO4189 nucleotide sequence is highly similar to SCO4190 (*devA*) encoding for a GntR-family transcriptional regulator required for morphological development (Hoskisson *et al.*, 2006). The high similarity is distributed along two stretches 23 and 26 nucleotides long (80% and 86% identity respectively); this result suggests a very likely annealing between the two transcripts. Therefore, SCO4189 might transcribe a ncRNA that could act on *devA* mRNA, decreasing its level. Strengthening this hypothesis, the differential expression profile, between the wild type and the *scbA* mutant, of SCO4189 and *devA* is anticorrelated (data not shown).

The detailed analysis of the differential expression at TP1 showed that many genes with housekeeping functions and some genes previously reported to affect antibiotic regulation were upregulated in M751 just before the onset of antibiotic production. Genes related to stress response, i.e. *scoF4*, *smpB*, *dnaJ* and *groEL2*, were upregulated in the mutant. Overexpression of *csp1* (homologue to *scoF4*) of *Streptomyces hygroscopicus* in *S. coelicolor* has been reported to induce overproduction of both Red and Act (Martinez-Costa *et al.*, 2003). Furthermore, several studies have been carried out to increase antibiotic production by shock stimuli, for example heat, cold, oxidative or osmotic stress (Hayes *et al.*, 1997; Mikulik *et al.*, 1999; Kim *et al.*, 2000; Sevcikova and Kormanec, 2004), and it is widely known that stress induces antibiotic production. Also the pleiotropic regulator *afsS* was found upregulated in M751. Overexpression of *afsS* in *S. coelicolor* (reviewed in Horinouchi, 2003) or its counterpart *afsR2* in *Streptomyces lividans* (Vogtli *et al.*, 1994) has been reported to enhance production of Act and Red. Among several genes encoding hypothetical proteins, SCO3893 is predicted to have a kinase domain and seems to be transcriptionally

coupled to *sigT* (SCO3892) and *rstA* (SCO3891), which have been described as extracytoplasmic functions (ECF) sigma factor and anti-sigma factor, both involved in differentiation of *S. coelicolor* (Mao *et al.*, 2009b). SCO2195 has been shown to be a target of the nitrogen regulator GlnR (Tiffert *et al.*, 2008). Because of the similarity with the system in *Rhizobium leguminosarum*, the authors speculate a possible interaction, and thus a regulatory function, of this small protein with the mRNA of *glnA*, encoding the main glutamine synthetase in *S. coelicolor*. Furthermore, *glnA* (SCO2198) was shown by our microarray analysis, to be downregulated in the mutant strain. These data are encouraging for the analysis of the other identified genes, in particular several hypothetical proteins, for their possible role in antibiotic production and morphological differentiation (see Table 1).

The present study portrays a strong link between primary and secondary metabolism and supports the idea that the SCB system has a possible role in coordinating cell metabolism in *S. coelicolor*. Furthermore, our data suggest a useful biotechnological approach for optimization of antibiotic production. We propose that manipulation of the  $\gamma$ -butyrolactones synthase can be a successful alternative for engineering antibiotic production in microbial systems.

## Experimental procedures

### Bacterial strains, plasmids and growth conditions

*Streptomyces* strains were manipulated as described previously (Kieser *et al.*, 2000). Supplemented minimal medium (SMM) (Takano *et al.*, 2001) was used for RNA isolation and antibiotic production determination. MS agar (Kieser *et al.*, 2000) was used to make spore suspensions.

### Cell culture, DNA and RNA isolation

For the microarray analysis and qRT-PCR, the same amount of *S. coelicolor* M145 and M751 spores was inoculated and grown in three biological replicates in SMM, and RNA was isolated at eight time points during the growth curve as previously reported (Takano *et al.*, 2001).

### cDNA preparation, labelling and microarray hybridization

cDNA preparation and labelling was performed as previously described (Takano *et al.*, 2005). Ten micrograms of labelled cDNA and 0.2  $\mu$ g of labelled genomic DNA (gDNA) were hybridized to PCR-based microarray slides from the University of Minnesota comprising 7629 genes. After 16 h of hybridization and three subsequent washing steps the arrays were scanned with an Affymetrix Array scanner 417.

### Analysis of microarray data

Raw data collection was performed using ImaGene (as described in Takano *et al.*, 2005). Further statistical and bio-

informatical analyses were performed using the R language (<http://www.r-project.org>) and packages from the Bioconductor project (limma, marray, multtest, hcluster). The data were normalized using print-tip loess normalization and between-array scale normalization available from the limma package. gDNA was used as common reference for all hybridizations. The relative fluorescence intensity of cDNA to gDNA ( $\log_2$  expression value) indicates the transcript abundance at each time point in each strain. Differentially expressed genes were calculated for the three replicates using Welch's two-sample *t*-test, the non-parametric Mann–Whitney test and the Rank Product test. The intersection of genes with *P*-value < 0.05 of the *t*-test and Mann–Whitney test was combined with genes that had a multiple-testing corrected *p* < 0.05 in the Rank Product test.

### cDNA synthesis and qRT-PCR

DNase treatment was performed on 10  $\mu$ g of each RNA sample with 10 U of DNase I (Roche), and the absence of chromosomal DNA contamination was checked by PCR using the primers hrdBroutfw/hrdBroutvr (Table S2). To synthesize cDNA, 2  $\mu$ g of RNA was annealed to 1.8  $\mu$ g of random hexamers (Invitrogen) and incubated at 70°C for 10 min; to this 5 $\times$  First-Strand Buffer, DTT to final 10 mM, dNTPs to final 1 mM and 100 U of Superscript II (Invitrogen) were added in a final volume of 30  $\mu$ l and the mixture was incubated at 25°C for 10 min, 37°C for 5 min and 42°C for 100 min. qRT-PCR experiments were performed using the ABI Prism 7500FAST system (Applied Biosystems). Each 25  $\mu$ l reaction contained 12.5  $\mu$ l of SYBR Green PCR Master Mix (Applied Biosystems), 10 ng of cDNA (or water for the negative control), 10 pmol of each primer (listed in Table S2) and 9.5  $\mu$ l of water. The reaction parameters were as following: 95°C for 10 min, followed by 40 two-step amplification cycles consisting of 30 s denaturation at 95°C and 1 min of annealing and extension at 60°C. A final dissociation step was run to generate a melting curve and consequently verify the specificity of amplification products. Real-time PCR was monitored and analysed by the Sequence Detection System version 1.3 (Applied Biosystems), and relative expression levels were normalized to mRNA of the major vegetative sigma factor (*hrdB*). All samples were run in triplicate.

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## Supporting information

Additional Supporting Information may be found in the online version of this article:

**Fig. S1.** Expression profiles of the butanolide cluster. Expression profiles of *scbA* (SCO6266) (values from qRT-PCR analysis), *scbR* (SCO6265) (values from qRT-PCR analysis) and *scbC* (SCO6267) (values from microarray analysis) are plotted for M145 (dark line) and the  $\Delta scbA$  mutant (light line). The expression fold change for *scbA* and *scbR* is compared with the value in M145 at TP1, which is set as 1. Standard deviation for *scbA* and *scbR* qRT-PCR analysis at each time point is denoted in a thin light line.



**Table S1.** Differentially expressed genes from TP1.

**Table S2.** Primers used for qRT-PCR.

**Table S3.** Expression data. Expression values of individual genes in the clusters shown in Figs 3, 5, 6 and 7. The columns indicate the SCO number, the gene name (if available) and the normalized expression value from the microarray analysis at each time point in the wild-type M145 and the mutant M751.

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